

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : C07C 67/38, 69/24, B01J 31/24	A1	(11) International Publication Number: WO 96/19434 (43) International Publication Date: 27 June 1996 (27.06.96)
(21) International Application Number: PCT/GB95/03021 (22) International Filing Date: 22 December 1995 (22.12.95) (30) Priority Data: 9425911.6 22 December 1994 (22.12.94) GB (71) Applicant (for all designated States except US): IMPERIAL CHEMICAL INDUSTRIES PLC [GB/GB]; Imperial Chem- ical House, Millbank, London SW1P 3JF (GB). (72) Inventors; and (75) Inventors/Applicants (for US only): TOOZE, Robert, Paul [GB/GB]; 11 Station Road, Norton, Stockton, Cleveland TS20 1EA (GB). EASTHAM, Graham, Ronald [GB/GB]; 7 Heslop Drive, Darlington, Co Durham DL1 5TQ (GB). WHISTON, Keith [GB/GB]; 22 Ravensdale Road, West End, Darlington, Co Durham DL3 8EA (GB). WANG, Xiao, Lan [CN/GB]; 108 Lambton Road, Marton Grove, Middlesbrough, Cleveland TS4 2RG (GB). (74) Agents: MILLROSS, Christopher et al.; ICI Materials, Intel- lectual Property Dept., P.O. Box 90, Wilton, Middlesbrough, Cleveland TS90 8JE (GB).	(81) Designated States: AM, AU, BB, BG, BR, BY, CA, CN, CZ, FI, GE, HU, JP, KE, KG, KP, KR, KZ, LK, LT, LV, MD, MG, MN, MW, NO, NZ, PL, RO, RU, SD, SI, SK, TJ, TT, UA, US, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>	
(54) Title: PROCESS FOR THE CARBONYLATION OF ETHYLENE AND CATALYST SYSTEM FOR USE THEREIN		
(57) Abstract A catalyst system capable of catalysing the carbonylation of ethylene and a process which uses such a system wherein the catalyst system is obtainable by combining a metal of Group VIII, e.g. palladium, or a compound thereof and a bidentate phosphine, e.g. bis (di-t-butyl phosphino) - o - xylene.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	GB	United Kingdom	MR	Mauritania
AU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IE	Ireland	NZ	New Zealand
BJ	Benin	IT	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgyzstan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LI	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

Process for the carbonylation of ethylene and catalyst system for use therein

The invention relates to the carbonylation of ethylene using carbon monoxide in the presence of a catalyst system and to such a catalyst system.

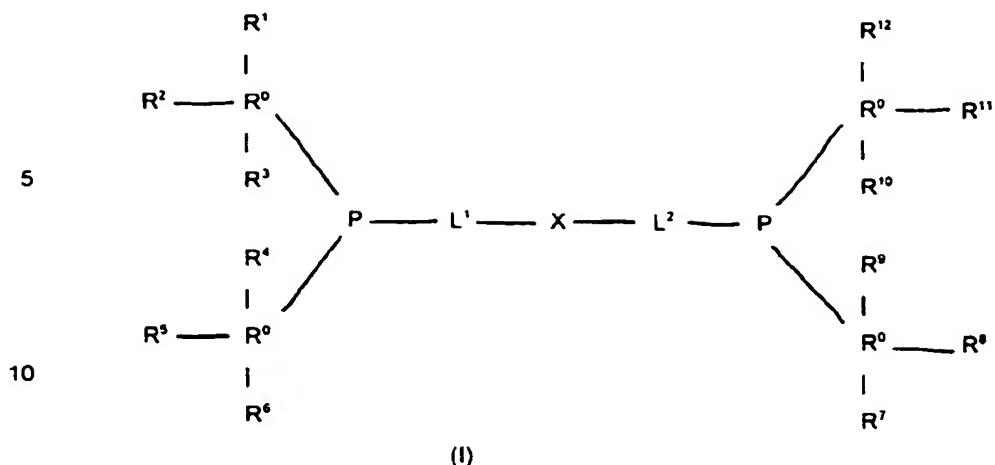
The carbonylation of ethylene using carbon monoxide in the presence of an alcohol 5 or water and a catalyst system comprising a Group VIII metal, e.g. palladium, and a phosphine ligand, e.g. an alkyl phosphine, cycloalkyl phosphine, aryl phosphine, pyridyl phosphine or bidentate phosphine, has been described in numerous European patents and patent applications, e.g. EP-A-0055875, EP-A-04489472, EP-A-0106379, EP-A-0235864, EP-A-0274795, EP-A-0499329, EP-A-0386833, EP-A-0441447, EP-A-0489472, 10 EP-A-0282142, EP-A-0227160, EP-A-0495547 and EP-A-0495548. In particular, EP-A-0227160, EP-A-0495547 and EP-A-0495548 disclose that bidentate phosphine ligands provide catalyst systems which enable higher reaction rates to be achieved.

The main problem with the previously disclosed catalyst systems is that, although relatively high reaction rates can be achieved, the palladium catalyst dies off quickly which 15 necessitates the frequent replenishment of the catalyst and hence results in a process which is industrially unattractive.

It has now been found that a particular group of bidentate phosphine compounds can provide remarkably stable catalysts which require little or no replenishment; that use of such bidentate catalysts leads to reaction rates which are significantly higher than those 20 previously disclosed; that little or no impurities are produced at high conversions.

Accordingly, the present invention provides a process for the carbonylation of ethylene which process comprises reacting ethylene with carbon monoxide in the presence of a source of hydroxyl groups and of a catalyst system, wherein the catalyst system is obtainable by combining:

- 25 (a) a metal of Group VIII or a compound thereof; and
 (b) a bidentate phosphine of general formula (I)



wherein

R^0 is a tertiary carbon atom

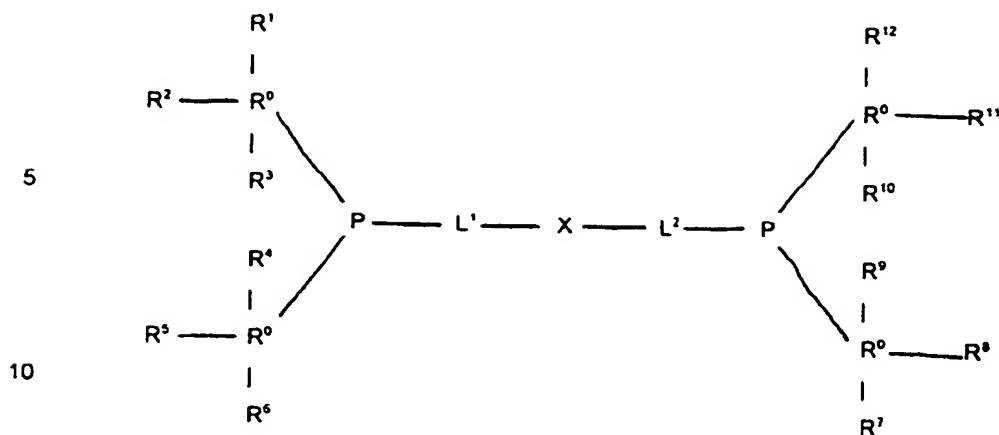
15 each of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} and R^{12} is independently a pendant optionally substituted organic group which carries a carbon atom through which the group is linked to the respective R^0 ;

each of L^1 and L^2 is independently a linking group selected from an optionally substituted lower alkylene chain connecting the respective phosphorus atom to the group X;
20 and

X is a bridging group comprising an optionally substituted aryl moiety to which the phosphorus atoms are linked on available adjacent carbon atoms.

In a second aspect, the present provides a catalyst system capable of catalysing the carbonylation of ethylene, which catalyst system is formed from

- 25
- (a) a metal of Group VIII or a compound thereof; and
 - (b) a bidentate phosphine of general formula (I)



wherein

R^0 is a tertiary carbon atom

15 each of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} and R^{12} is independently a pendant optionally substituted organic group which carries a carbon atom through which the group is linked to the respective R^0 ,

each of L^1 and L^2 is independently a linking group selected from an optionally substituted lower alkylene chain connecting the respective phosphorus atom to the group X;
20 and

X is a bridging group comprising an optionally substituted aryl moiety to which the phosphorus atoms are linked on available adjacent carbon atoms.

The pendant optionally substituted organic groups, R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} and R^{12} , may be independently selected from a wide range of components.
25 Preferably, the pendant groups are optionally substituted lower alkyl, e.g. C_{1-6} , and which may be branched or linear.

Particularly preferred is when the organic groups, R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} and R^{12} , when associated with their respective R^0 carbon atom form composite groups which are at least as sterically hindering as t-butyl. Steric hindrance in this context is
30 as discussed at page 14 et seq of "Homogeneous Transition Metal Catalysis - A Gentle Art", by C Masters, published by Chapman and Hall, 1981.

The linking groups, L^1 and L^2 , are independently selected from an optionally substituted, particularly lower alkyl, e.g. C_1 to C_4 , substituted, lower alkylene, e.g. C_1 to C_4 chain. Especially preferred is when both L^1 and L^2 are methylene.

35 The bridging group X is an aryl moiety, e.g. a phenyl group, which may be optionally

substituted, provided that the two phosphorus atoms are linked to adjacent carbon atoms, e.g. at the 1 and 2 positions on the phenyl group. Optional substitution of the aryl moiety may be by other organic groups, e.g. alkyl, particularly C_{1-8} , aryl, alkoxy, carbalkoxy, halo, nitro, trihalomethyl and cyano. Furthermore, the aryl moiety may be a fused polycyclic group, e.g. 5 naphthalene, biphenylene or indene.

Examples of suitable bidentate ligands are bis (di-t-butyl phosphino) - o - xylene (also known as 1,2 bis (di-t-butylphosphinomethyl) benzene), bis (di-t-neopentyl phosphino) - o - xylene and bis 1, 2 (di-t-butyl phosphino) naphthalene. Additionally, the bidentate phosphine may be bonded to a suitable polymeric substrate via at least one of the bridging 10 group X, the linking group L^1 or the linking group L^2 , e.g. bis (di-t-butyl phosphino) - o - xylene may be bonded via the xylene group to polystyrene to give an immobile heterogeneous catalyst.

The amount of bidentate ligand used can vary within wide limits. Preferably, the bidentate ligand is present in an amount such that the ratio of the number of moles of the 15 bidentate ligand present to the number of moles of the Group VIII metal present is from 1 to 50, e.g. 1 to 10 and particularly from 1 to 5 mol per mol.

The carbon monoxide may be used in the presence of other gases which are inert in the reaction. Examples of such gases include hydrogen, nitrogen, carbon dioxide and the noble gases such as argon.

20 The process of the present invention is preferably carried out at a temperature from 20 to 250°C, in particular from 40 to 150°C and especially from 70 to 120°C.

The process may be conducted under a total pressure of from 1×10^5 to 100×10^5 N.m⁻² and in particular from 5×10^5 to 50×10^5 N.m⁻².

Suitable Group VIII metals include cobalt, nickel, palladium, rhodium and platinum. 25 Particularly preferred is palladium. Suitable compounds of such Group VIII metals include salts of such metals with, or compounds comprising weakly co-ordinated anions derived from, nitric acid; sulphuric acid; lower alkanolic (up to C_{12}) acids such as acetic acid and propionic acid; sulphonic acids such as methane sulphonic acid, chlorosulphonic acid, fluorosulphonic acid, trifluoro methane sulphonic acid, benzene sulphonic acid, naphthalene sulphonic acid, 30 toluene sulphonic acid, e.g. p-toluene sulphonic acid, t-butyl sulphonic acid, and 2-hydroxypropane sulphonic acid; sulphonated ion exchange resins; perhalic acid such as perchloric acid; perfluorinated carboxylic acid such as trichloroacetic acid and trifluoroacetic acid; orthophosphonic acid; phosphonic acid such as benzene phosphonic acid; and acids derived from interactions between Lewis acids and Broensted acids. Other sources which 35 may provide suitable anions include the tetraphenyl borate derivatives. Additionally, zero

valent palladium compounds with labile ligands, e.g. tri (dibenzylideneacetone) dipalladium, may be used.

The catalyst system of the present invention may be used homogeneously or heterogeneously. Preferably the catalyst system is used homogeneously.

5 The catalyst system of the present invention is preferably constituted in the liquid phase which may be formed by one or more of the reactants or by the use of a suitable solvent.

Suitable solvents that may be used in conjunction with the catalyst system include one or more aprotic solvents such as ethers, e.g. diethyl ether, dimethyl ether, dimethyl ether
10 of diethylene glycol, anisole and diphenyl ether; aromatic compounds, including halo variants of such compounds, e.g. benzene, toluene, ethyl benzene, o-xylene, m-xylene, p-xylene, chlorobenzene, o-dichlorobenzene, m-dichlorobenzene, and p-dichlorobenzene; alkanes, including halo variants of such compounds, e.g. hexane, heptane, 2, 2, 3- trimethylpentane, methylene chloride and carbon tetrachloride; nitriles, e.g. benzonitrile and acetonitrile; esters,
15 e.g. methyl benzoate, methyl acetate and dimethyl phthalate; sulphones, e.g. diethyl sulphone and tetrahydrothiophene 1, 1- dioxide; amides, including halo variants of such compounds, e.g. dimethyl formamide and N-methyl pyrrolidone.

The catalyst system of the present invention is particularly suited to the carbonylation of ethylene. Surprisingly, propene has been found to be difficult to carbonylate
20 to the extent that the present catalyst system may be viewed as not being able to carbonylate propene.

The end product of the reaction is determined at least in part by the source of hydroxyl groups that is used. The use of water gives rise to the corresponding carboxylic acid whereas the use of an alcohol leads to the corresponding ester. Suitable alcohols include
25 C₁₋₃₀ alcohols, optionally substituted with one or more substituents such as halogen atoms, cyano, carbonyl, alkoxy or aryl groups. Suitable alcohols include methanol, ethanol, propanol, 2-propanol, 2-butanol, 1-butyl alcohol and chlorocapryl alcohol. Particularly useful are methanol and ethanol.

The molar ratio of the amount of ethylene used in the reaction to the amount of
30 hydroxyl providing compound is not critical and may vary between wide limits, e.g. from 0.001:1 to 100:1 mol/mol.

The product of the reaction may be separated from the other components by any suitable means. However, it is an advantage of the present catalyst system that significantly
fewer by-products are formed thereby reducing the need for further purification after the initial
35 separation of the product as may be evidenced by the generally significantly higher

selectivity. A further advantage is that the other components which contain the catalyst system which may be recycled and/or reused in further reactions with minimal supplementation of fresh catalyst.

The following Examples further illustrated the present invention.

5 Example 1

In this example methyl propionate was prepared from carbon monoxide and ethylene using methanol as the hydroxyl group source.

A mechanically stirred autoclave of 2 litre capacity was evacuated of air and then charged with a catalyst system consisting of

10	methanol	300 cm ³
	palladium acetate	0.1 mmol
	bidentate*	0.3 mmol
	methane sulphonic acid	0.24 mmol

* - bis (di-t-butylphosphino) - o - xylene

15 Carbon monoxide and ethylene on an equimolar basis was introduced until a pressure of $30 \times 10^5 \text{ N.m}^{-2}$ was reached. The temperature of the reactor was raised to and maintained at 100°C. As the reaction proceeded sufficient additional methanol was introduced to compensate for that which had been consumed and additional carbon monoxide and ethylene was added (on an equimolar basis) to maintain the pressure. No palladium
20 precipitation was observed.

Example 2 - Comparative

Example 1 was repeated except that the propane analogue of the o-xylene bidentate was used. Precipitation of palladium, with consequent deactivation of the catalyst, was observed within two hours.

25 The reaction rate (expressed as mole ethylene per mole of palladium per hour), selectivity to methyl propionate (% by gas chromatography) and turn over number based on phosphine (expressed as mole methyl propionate per mole of phosphine) for each of the catalyst systems described in Examples 1 and 2 is shown in Table 1 below.

TABLE 1

PHOSPHINE LIGAND	REACTION RATE	SELECTIVITY	TURN OVER NUMBER
bis (di-t-butyl phosphino)-o-xylene	40000	99.95	>50000
bis (di-t-butyl phosphino) - propane	15000	98.00	1700

It can thus be seen that the catalyst system of the present invention is more stable, more reactive and more selective than the conventional bidentate system.

Example 3

- 5 The catalyst system from Example 1 was extracted at the end of the procedure and reused with fresh methanol and methane sulphonic acid. The activity of the reused catalyst was the same as that of the original. The catalyst system of Example 2 was unable to be reused due to the precipitation of the palladium.

Example 4

- 10 Example 1 was repeated using the 4-nitro substituted analogue of the bidentate ligand which was prepared via the phosphonium salt produced from the reaction of the appropriate secondary phosphine with the corresponding aromatic dihalide.

Example 5

Example 4 was repeated using the 4-methoxy substituted analogue.

- 15 The results of Example 4 and 5 were as follows:

PHOSPHINE LIGAND	REACTION RATE	SELECTIVITY	TURN OVER NUMBER
4-nitro	36000	99.9	>25000
4-methoxy	37000	99.9	>25000

Example 6

Example 1 was repeated except that an alternative source of palladium was used. The catalyst was prepared by mixing the bidentate phosphine, tris(dibenzylideneacetone) dipalladium known as dba, and sulphonic acid in a molar ratio of 2:1:3. The phosphine and

dba were mixed together prior to the addition of the acid.

The reaction was conducted using an equimolar mixture of methanol and methyl propionate at a total pressure of 15 barg (using a 1:1 mixture of carbon monoxide and ethylene). The reaction temperature was 80°C.

5 Example 7

Example 6 was repeated except that in the ligand one of the t-butyl groups on each phosphorous atom was replaced by cyclohexyl groups.

Example 8

Example 6 was repeated except that in the ligand the t-butyl groups were replaced
10 by cyclohexyl groups.

Example 9

Example 6 was repeated except that in the ligand the t-butyl groups were replaced
by isopropyl groups.

Example 10

15 Example 6 was repeated except that in the ligand the t-butyl groups were replaced
by phenyl groups.

Example 11

Example 6 was repeated except that the linking methylene groups were both
replaced by oxygen atoms. The phosphinite was synthesised from an aromatic diol and
20 chloro di(t-butyl) phosphine.

Example 12

Example 11 was repeated except that the 4 t-butyl substituted analogue of the
ligand was used.

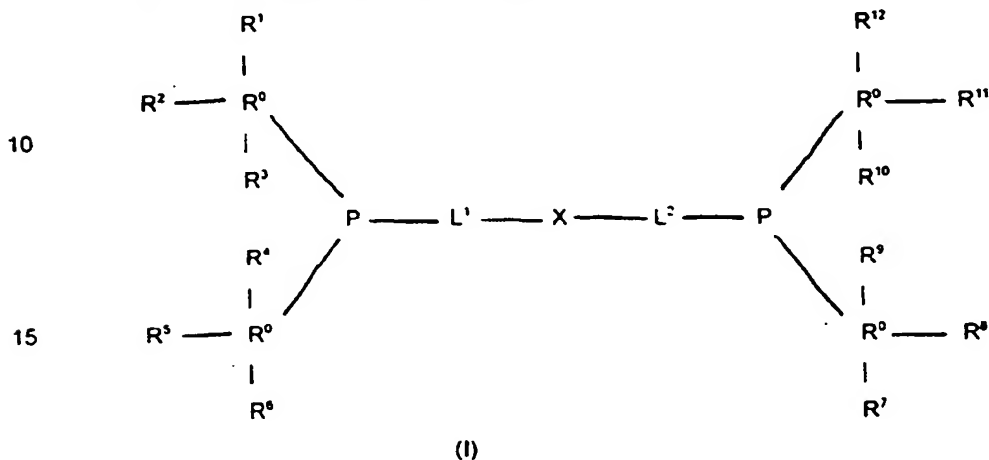
The results of Examples 6 to 12 were as follows:

PHOSPHINE LIGAND	REACTION RATE	SELECTIVITY	TURN OVER NUMBER
Example 6	12000	99.95	>250000
Example 7	500	30	>1500
Example 8	200	25	>600
Example 9	200	20	500
Example 10	400	20	>1200
Example 11	100	30	<300
Example 12	100	30	<300

Claims

1. A process for the carbonylation of ethylene which process comprises reacting ethylene with carbon monoxide in the presence of a source of hydroxyl groups and of a catalyst system, wherein the catalyst system is obtainable by combining:

- 5 (a) a metal of Group VIII or a compound thereof; and
 (b) a bidentate phosphine of general formula (I)



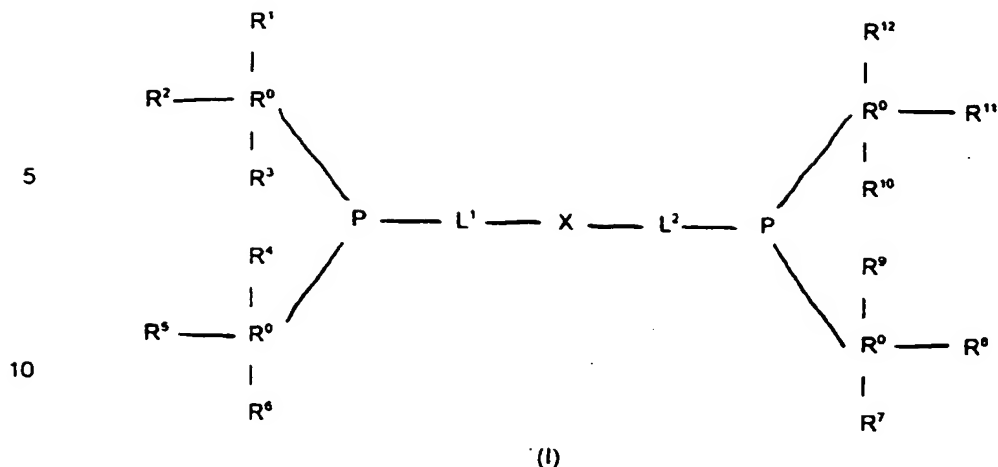
wherein

- 20 R^0 is a tertiary carbon atom
 each of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} and R^{12} is independently a pendant optionally substituted organic group which carries a carbon atom through which the group is linked to the respective R^0 ;
 each of L^1 and L^2 is independently a linking group selected from an optionally substituted lower alkylene chain connecting the respective phosphorus atom to the group X;
 25 and

X is a bridging group comprising an optionally substituted aryl moiety to which the phosphorus atoms are linked on available adjacent carbon atoms.

2. A catalyst system capable of catalysing the carbonylation of an ethylene, which
 30 catalyst system is formed from

- (a) a metal of Group VIII or a compound thereof; and
 (b) a bidentate phosphine of general formula (I)



wherein

R^0 is a tertiary carbon atom

15 each of R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} and R^{12} is independently a pendant optionally substituted organic group which carries a carbon atom through which the group is linked to the respective R^0 ;

each of L^1 and L^2 is independently a linking group selected from an optionally substituted lower alkylene chain connecting the respective phosphorus atom to the group X;
20 and

X is a bridging group comprising an optionally substituted aryl moiety to which the phosphorus atoms are linked on available adjacent carbon atoms.

3. A process or catalyst system as claimed in claim 1 or claim 2 wherein the pendant optionally substituted organic groups, R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} and R^{12} , are
25 independently selected from optionally substituted lower alkyl.

4. A process or catalyst system as claimed in any one of the preceding claims wherein the organic groups, R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} and R^{12} , are associated with their respective R^0 carbon atom so as to form composite groups which are at least as sterically hindering as t-butyl.

30 5. A process or catalyst system as claimed in any one of the preceding claims wherein both L^1 and L^2 are methylene.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 95/03021

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07C67/38 C07C69/24 B01J31/24

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07C B01J

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP,A,0 495 548 (SHELL INTERNATIONALE RESEARCH MAATSCHAPPIJ B.V.) 22 July 1992 cited in the application see page 2, line 39 - line 55 see page 11 - page 12; claims ---	1
X	EP,A,0 375 573 (EASTMAN KODAK COMPANY) 27 June 1990 see page 3, line 19 - line 58 see page 7, line 8 - line 15 see page 8 - page 9; example 1 see page 19 - page 20; claims 20-22 -----	2,5

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- *&* document member of the same patent family

Date of the actual completion of the international search

20 March 1996

Date of mailing of the international search report

02.04.96

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax (+31-70) 340-3016

Authorized officer

Kinzinger, J

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No
PCT/GB 95/03021

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP-A-495548	22-07-92	CA-A- 2059236	16-07-92
		DE-D- 69204691	19-10-95
		JP-A- 5058949	09-03-93
		CA-A- 2059233	16-07-92
		CN-A- 1063277	05-08-92
		EP-A- 0495547	22-07-92
		JP-A- 4334340	20-11-92

EP-A-375573	27-06-90	US-A- 4960949	02-10-90
		CA-A- 2004707	22-06-90
		DE-T- 68905452	21-10-93
		EP-A- 0449966	09-10-91
		JP-T- 4502463	07-05-92
		WO-A- 9006810	28-06-90
		US-A- 5004823	02-04-91

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

☐ **BLACK BORDERS**

☐ **IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**

☒ **FADED TEXT OR DRAWING**

☐ **BLURRED OR ILLEGIBLE TEXT OR DRAWING**

☐ **SKEWED/SLANTED IMAGES**

☐ **COLOR OR BLACK AND WHITE PHOTOGRAPHS**

☐ **GRAY SCALE DOCUMENTS**

☐ **LINES OR MARKS ON ORIGINAL DOCUMENT**

☐ **REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**

☐ **OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.